



IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re application of

Confirmation No. 9517

Tsuyoshi SUZUKI et al.

Attorney Docket No. 2004 1909A

Serial No. 10/516,360

Group Art Unit 1625

Filed March 4, 2005

Examiner Taofiq A. Solola

PREVENTIVES AND/OR REMEDIES FOR SUBJECTS WITH THE EXPRESSION OR

ACTIVATION OF Her2 AND/OR EGFR

Mail Stop: AMENDMENT

SUPPLEMENTAL REPLY

Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

Sir:

Attached to this paper is a Declaration by Tsuyoshi Suzuki, executed June 27, 2008. This Declaration is the original, a copy of this Declaration was submitted with our response to the Office Action of April 3, 2008. Please replace the copy Declaration with this attached Declaration.

Respectfully submitted,

Tsuyoshi SUZUKI et al.

By: /hlallan 91/

William R. Schmidt, II Registration No. 58,327 Attorney for Applicants

WRS/lc Washington, D.C. 20006-1021 Telephone (202) 721-8200 Facsimile (202) 721-8250 July 10, 2008

THE COMMISSIONER IS AUTHORIZED TO CHARGE ANY DEFICIENCY IN THE FEES FOR THIS PAPER TO DEPOSIT ACCOUNT NO. 23-0975



IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re application of : Confirmation No. 9517

Tsuyoshi SUZUKI et al. : Attorney Docket No. 2004 1909A

Serial No. 10/516,360 : Group Art Unit 1625

Filed March 4, 2005 : Examiner Taofiq A. Solola

PREVENTIVES AND/OR REMEDIES FOR SUBJECTS WITH THE EXPRESSION OR

ACTIVATION OF Her2 AND/OR EGFR : Mail Stop: AMENDMENT

DECLARATION UNDER 37 C.F.R. § 1.132

Assistant Commissioner for Patents, Washington, D.C.

Sir:

- I, Tsuyoshi Suzuki, the undersigned, a citizen of Japan, residing at 2-30-40 Sumiyoshi, Fuchu, Tokyo do hereby declare:
 - 1. That I am an inventor of the above-identified application.
- 2. That I graduated from Tohoku University, graduate school of pharmaceutical science with a Master's degree in 1992.
- 3. I am an employee of Mitsubishi Tanabe Pharma Corporation. I have joined the company and been doing research of anticancer agents since 1992.
- 4. Relevant Publication: Suzuki T, Fujii A, Ohya J et al. Pharmacological characterization of MP-412 (AV-412), a dual epidermal growth factor receptor and ErbB2 tyrosine kinase inhibitor. Cancer Science 2007; 98: 1977-1984.
- 5. That in order to show enablement under 35 U.S.C. § 112 of Claims 19-24 of the above-identified application, I have under my control and direction conducted the following experiments. The particulars and results of the experiments are set forth hereinbelow.

EXPERIMENTAL DATA

- 1. Object: In vivo antitumor effects
- 2. Method: Antitumor effects of compound A were evaluated by using various human cancer cell lines which express Her2 and/or EGFR. Human lung cancer NCI-H1975 (ATCC No. CRL-5908), prostate cancer DU145 (ATCC No. HTB-81) and ovarian cancer SKOV-3 (ATCC No. HTB-77) were purchased from ATCC. Human esophageal cancer TE-8 was obtained from the Cell Resource Center for Biomedical Research (Tohoku University). Human Breast cancer KPL-4 was gift from Dr. Junichi Kurebayashi (Kawasaki Medical University). The chemical name and the structure of compound A are as follows.

N-{4-[(3-Chloro-4-fluorophenyl)amino]-7-[3-methyl-3-(4-methyl1-piperazinyl)-1-butynyl]-6-quinazolinyl}acrylamide

$$\begin{array}{c|c} H_3C & CH_3 \\ N & & \\ H_2C & & \\ \end{array}$$

3. Results: The results are shown in the following Tables.

(Table x1)

Antitumor effects on human lung cancer NCI-H1975 (both EGFR, Her2 positive)				
Pharmaceutical agent	Dose mg/kg	Average relative tumor growth rate	% control	
Control	-	5.5	100	
Compound A · 2TsOH	75	3.1	47	
Compound A· 2TsOH	150	0.9	-2	

(Table x2)

Antitumor effects of	on human breast ca	ancer KPL-4 (both EGF	R, Her2 positive)
Pharmaceutical	Dose mg/kg	Average relative	% control
agent		tumor growth rate	
Control	-	4.8	100
Compound A.	100	1.5	14
2TsOH			

(Table x3)

Antitumor effects or	human prostate c	ancer DU145 (both EGI	R, Her2 positive)
Pharmaceutical agent	Dose mg/kg	Average relative tumor growth rate	% control
Control	_	3.3	100
Compound A · 2TsOH	100	1.3	11

(Table x4)

Antitumor effects or	n human ovarian ca	ancer SKOV-3 (both EGI	FR, Her2 positive)
Pharmaceutical agent	Dose mg/kg	Average relative tumor growth rate	% control
Control	-	9.6	100
Compound A· 2TsOH	100	6.1	60

(Table x5)

Antitumor effect	•	ageal cancer TE-8 (both sitive)	n EGFR, Her2
Pharmaceutical agent	Dose mg/kg	Average relative tumor growth rate	% control
Control	<u>.</u> .	5.6	100
Compound A · 2TsOH	10	0.34	-14

(Table x6)

Antitumor effect		eatic cancer HPAC (both sitive)	n EGFR, Her2
Pharmaceutical agent	Dose mg/kg	Average relative tumor growth rate	% control
Control	-	6.6	100
Compound A · 2TsOH	100	2.9	33

(Table x7)

Antitumor effects	on human cervica	cancer ME-180 (EGFR	R positive, Her2
	neç	gative)	
Pharmaceutical agent	Dose mg/kg	Average relative tumor growth rate	% control
Control	-	9.6	100
Compound A · 2TsOH	100	6.1	60

(Table x8)

Antitumor effects of		al cancer LS174T (EGFF sitive)	R negative, Her2
Pharmaceutical agent	Dose mg/kg	Average relative tumor growth rate	% control
Control	-	. 13.9	100
Compound A· 2TsOH	100	8.2	56

(Table x9)

Antitumor effects on human lung cancer NCI-H520 (both EGFR, Her2 negative)				
Pharmaceutical	Dose mg/kg	Average relative	% control	
agent		tumor growth rate		
Control	-	7.4	100	
Compound A	100	6.5	86	
2TsOH				

5. Conclusion: From the results shown above in the tables, it is my expert opinion and belief that compound A suppresses the growth of various human cancer types which possess Her2 and/or EGFR. Thus, it is my expert opinion and belief that the claimed invention is enabled.

I further declare that all statements made herein of my own knowledge are true and all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of this application or any patent issuing thereon.

Date: June 27, 2008

(Signature of Declarant)